

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Centers for Medicare & Medicaid Services** 

[Document Identifier CMS-10701, CMS-10191 and CMS-10142]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, HHS

**ACTION:** Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the *Federal Register* concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments on the collection(s) of information must be received by the OMB desk officer by **[INSERT DATE 30 DAYS AFTER THE DATE OF PUBLICATION IN THE FEDERAL REGISTER]**.

**ADDRESSES:** When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions:

OMB, Office of Information and Regulatory Affairs

Attention: CMS Desk Officer

Fax Number: (202) 395-5806 OR

E-mail: *OIRA\_submission@omb.eop.gov* 

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web Site address at Web Site address at

https://www.cms.gov/Regulations-and-

Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html

1. E-mail your request, including your address, phone number, OMB number, and CMS

document identifier, to Paperwork@cms.hhs.gov.

2. Call the Reports Clearance Office at (410) 786-1326.

**FOR FURTHER INFORMATION CONTACT:** William Parham at (410) 786-4669.

**SUPPLEMENTARY INFORMATION:** Under the Paperwork Reduction Act of 1995 (PRA)

(44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management

and Budget (OMB) for each collection of information they conduct or sponsor. The term

"collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes

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agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the *Federal Register* concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

OMB control number); *Title of Information Collection:* Medicare Beneficiary Experiences with Care Survey System; *Use:* The MBECS system is designed to conduct 1-2 surveys per year on priority groups of interest, thereby allowing CMS OMH to respond quickly to the data needs of stakeholders with interests in these underrepresented groups. Data collected through the MBECS system will be used to better understand—and thus serve the needs of—Medicare beneficiaries in minority populations. The core questionnaire will collect information on communication with medical professionals, coordination of health care, experiences getting needed health care, experiences with personal doctors and specialists, and key demographics.

Data will be compared to benchmarks from the FFS CAHPS, MA CAHPS, and NAM CAHPS surveys. The population-specific questionnaire module will collect information about issues most relevant for particular minority groups; population-specific modules will be described in individual information collection requests. These data will be compared to benchmarks from the relevant CAHPS source surveys when available.

Collection of these data from people who have been identified through CMS administrative data and administrative flags as part of specific minority populations will also serve as a critical validation step of this method for identifying difficult-to-study populations, thus making it easier to study beneficiaries in these groups in the future. *Form Number*: CMS-10701 (OMB control number: 0938-NEW); *Frequency*: Yearly; *Affected Public*: State, Local, or Tribal Governments; *Number of Respondents*: 10,000; *Total Annual Responses*: 10,000; *Total Annual Hours*: 3,333. (For policy questions regarding this collection contact Luis Perez at 410-786-8557.)

2. Type of Information Collection Request: Revision with change of a currently approved collection; Title of Information Collection: Medicare Parts C and D Program Audit Protocols and Data Requests; Use: Under the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 and implementing regulations at 42 CFR Parts 422 and 423, Medicare Part D plan sponsors and Medicare Advantage organizations are required to comply with all Medicare Parts C and D program requirements. CMS' annual audit plan ensures that we evaluate sponsoring organizations' compliance with these requirements. CMS program audits focus on high-risk areas that have the greatest potential for beneficiary harm. As such, CMS has developed several audit protocols that are included within the program area data request documents and that are posted to the CMS website each year for use by sponsoring organizations to prepare for their audit. As part of a robust audit process, CMS also requires sponsoring organizations who have been audited and found to have deficiencies to undergo a validation audit to ensure correction. The validation audit utilizes the same audit protocols, but only tests

the elements where deficiencies were found, as opposed to re-administering the entire audit.

Currently CMS utilizes the following 5 protocols to audit sponsoring organization performance: Part D Formulary and Benefit Administration (FA); Part D Coverage Determinations, Appeals, and Grievances (CDAG); Part C Organization Determinations, Appeals, and Grievances (ODAG); Special Needs Model of Care (SNP-MOC) (only administered on organizations who operate SNPs); and, Compliance Program Effectiveness (CPE). The data collected is detailed in each of these protocols and the exact fields are located in the record layouts, at the end of each protocol. In addition, this collection request includes a preaudit issue summary, three CPE questionnaires, one CPE organizational structure presentation template, one FA impact analysis template, two CDAG impact analysis templates, four ODAG impact analysis templates, three SNP-MOC impact analysis templates, and a SNP-MOC questionnaire.

The information gathered during this audit will be used by the Medicare Parts C and D Oversight and Enforcement Group (MOEG) within the Center for Medicare (CM) and CMS Regional Offices to assess sponsoring organizations' compliance with Medicare program requirements. If outliers or other data anomalies are detected, Regional Offices will work in collaboration with MOEG and other divisions within CMS for follow-up and resolution. Additionally, MA and Part D organizations will receive the audit results and will be required to implement corrective action to correct any identified deficiencies. *Form Number*: CMS-10191 (OMB control number: 0938-1000); *Frequency*: Yearly; *Affected Public*: State, Local, or Tribal Governments; *Number of Respondents*: 201; *Total Annual Responses*: 207;

*Total Annual Hours:* 39,456. (For policy questions regarding this collection contact Brenda Hudson at 443-743-9299.)

3. Type of Information Collection Request: Revision with change of a currently approved collection; Title of Information Collection: Bid Pricing Tool (BPT) for Medicare Advantage (MA) Plans and Prescription Drug Plans (PDP); Use: Under the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (MMA), and implementing regulations at 42 CFR, Medicare Advantage organizations (MAOs) and Prescription Drug Plans (PDPs) are required to submit an actuarial pricing "bid" for each plan offered to Medicare beneficiaries for approval by the Centers for Medicare & Medicaid Services (CMS).

Medicare Advantage organizations (MAO) and Prescription Drug Plans (PDP) are required to submit an actuarial pricing "bid" for each plan offered to Medicare beneficiaries for approval by CMS. The MAOs and PDPs use the Bid Pricing Tool (BPT) software to develop their actuarial pricing bid. The competitive bidding process defined by the "The Medicare Prescription Drug, Improvement, and Modernization Act" (MMA) applies to both the MA and Part D programs. It is an annual process that encompasses the release of the MA rate book in April, the bid's that plans submit to CMS in June, and the release of the Part D and RPPO benchmarks, which typically occurs in August. *Form Number*: CMS-10142 (OMB control number: 0938-0944); *Frequency*: Yearly; *Affected Public*: State, Local, or Tribal Governments; *Number of Respondents*: 555; *Total Annual Responses*: 4995; *Total Annual Hours*: 149,850. (For policy questions regarding this collection contact Rachel Shevland at 410–786-3026.)

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Director,

Paperwork Reduction Staff,

Office of Strategic Operations and Regulatory Affairs.

4120-01-U-P

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